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Amendments to the Claims:

- 1. (Cancelled)
- 2. (Currently Amended) The use method as claimed in claim 9 1, wherein said linear n-alkanols are linear, possibly branched, hydrocarbon-chain n-alkanols in which have the OH group is in the 1-position (primary alcohol) or in the 2-position (secondary alcohol).
 - 3. (Cancelled)
- 4. (Currently Amended) The use method as claimed in claim 9 1, wherein said mammal has pathologies related to dysfunction of said CFTR are selected from the group consisting of cystic fibrosis, atypical cystic fibrosis, and obstructions of the bronchial tracts or of the digestive tracts.
- 5. (Currently Amended) The use method as claimed in claim 9 1, wherein said nalkanols are provided in a form suitable for intranasal or buccal administration.
- 6. (Currently Amended) The use method as claimed in claim 5, wherein said nalkanols are provided in a liquid form, for administration in the form of an aerosol or in the form of a nebulized material.
- 7. (Currently Amended) The use method as claimed in claim 6, wherein said nalkanols are combined with at least one pharmaceutically acceptable carrier appropriate for said intranasal or buccal administration.

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8. (Currently Amended) The use method as claimed in claim 9 1, wherein said nalkanols are administered at a concentration of between 0.001% and 0.1% (v/v), corresponding to a value of between 10 and 1000 ppm (parts per million), namely i.e. from 10 mg/kg to 1 g/kg.

9. (New) A method for partially or fully activating cystic fibrosis transmembrane conductance regulator channels (CFTR) in cell membranes of a mammal in need of such treatment comprising administering to said mammal at least one linear n-alkanol selected from the group consisting of C^6 - C_{10} and mixtures thereof in an amount sufficient to generate in the vicinity of said cell membranes a concentration of said n-alkanol sufficient to partially or fully open said CFTR in said cell membranes.